#### **IN THE CLAIMS:**

Please amend the following claims:

- 1 1. (Currently Amended) A process for the wet fractionation of cereal bran components,
- 2 [characterized in that] wherein bran is first subjected to a combination of enzymatic treatment
- 3 with enzymes of the group starch- and phytate-hydrolysing enzymes, and aqueous wet milling,
- 4 followed by an optional step of enzyme inactivation by wet heat treatment, and a subsequent step
- 5 whereby the insoluble phase containing a cleaned bran consisting of both pericarp and aleurone
- 6 fractions are separated by centrifugal forces into an aqueous phase containing a germ-rich
- 7 fraction and a further aqueous phase containing residual endosperm components, and that the
- 8 proteins contained in the endosperm-rich fraction are concentrated.
- 1 2. (Currently Amended) A process according to claim 1, wherein cereal brans are the fibrous-
- 2 residue resulting from a primary grain milling, i.e. after the separation of the endosperm fraction,
- 3 of wheat, rice, barley, oat, rye and triticale, and having variable chemical compositions, presence
- 4 of anti-nutritive factors, and presence of various anatomical fractions, i.e. pericarp, germ, and
- 5 residual endosperm.
- 1 3. (Currently Amended) A process according to claim 1, wherein the enzymatic treatment is
- 2 accomplished using a starch degrading enzyme of the group of amylases and amyloglucosidases.
- 4. (Currently Amended) A process according to [elaims 1-3] claim 1, wherein a further

- 2 enzymatic treatment is carried out using at least one non-starch degrading polysaccharidase in
- 3 the form of cellulases, hemicellulases mainly xylanases, beta-glucanases, and pectinases, and/or
- 4 phytases.
- 5. (Currently Amended) A process for the wet fractionation of cereal bran substantially free of
- 2 soluble compounds produced according to [elaim 1-3] claim 1, wherein such cleaned bran is
- 3 subjected to a combination of enzymatic treatment with specific enzymes of the group xylanase
- 4 and/or beta-glucanase under strictly controlled hydrolysis conditions, and intermittent wet
- 5 milling, followed by an optional step of enzyme inactivation by wet heat treatment.
- 6. (Currently Amended) A process according to claim 5, wherein the inactivated hydrolysate is
- 2 then fractionated by centrifugal forces into an insoluble phase containing primarily cellulose,
- 3 'lignin, less accessible hemicellulose, residual aleurone cells and cell wall bound proteins, and an
- 4 aqueous phase containing soluble hemicellulose, oligosaccharides, sugars and proteins, and that
- 5 the aqueous phase is further separated by centrifugal force into protein-rich fraction and a
- 6 carbohydrate-rich fraction, and that the carbohydrate-rich fraction is further separated by size
- 7 exclusion technique into a hemicellulose-rich fraction (medium molecular size fraction) and an
- 8 oligosaccharide-rich fraction (small molecular size fraction).
- 7. (Currently Amended) A process according to [claims 5-6] claim 5, wherein cereal bran
- 2 substantially free of both in water or less polar solvents soluble compounds are derived from
- 3 wheat, rice, barley, oat, rye or triticale.

- 8. (Currently Amended) A process according to [elaims 1 and 5-7] claim 1, wherein the
- 2 combination of intermittent wet milling with enzymatic treatment is arranged to increase the rate
- 3 of enzymatic hydrolysis of the substrate thereby improving the overall hydrolysis performance
- 4 and the subsequent separation of the various fractions by density/solubility and molecular size.
- 9. (Currently Amended) A process according to [elaims 5-8] claim 5, wherein the enzymatic
- 2 treatment is carried out using at least one non-starch degradable polysaccharidase in the form of
- 3 cellulases, hemicellulases mainly xylanases, beta-glucanases, and pectinases, and optionally
- 4 phytases.
- 1 10. (Currently Amended) A process according to claim 9, wherein the enzymatic treatment is
- 2 accomplished by using xylanases with high beta 1-4- xylanase (pentosanase) and/or beta-
- 3 glucanase activity.
- 1 11. (Currently Amended) A protein fraction derived substantially from the germ and produced
- 2 according to [elaims 1-4] claim 1, wherein the said fraction contains at least 35% protein and
- 3 10% oil on dry matter basis and exhibits a high emulsifying capacity and an increased shelf life
- 4 with regards to resistance to oxidation compared to the original bran, and that the said fraction
- 5 contains less than 5% fibre.
- 1 12. (Currently Amended) A protein fraction derived substantially from the residual endosperm
- and produced according to [elaims 1-4] claim 1, wherein the said fraction contains at least 25%

- 3 protein and 10% sugar and less than 3% oil and 3% fibre, and at least 25% soluble high-
- 4 molecular weight non-starch polysaccharides of the groups beta-glucans for barley and oat and
- 5 arabinoxylans for wheat, rice, rye and triticale.
- 1 13. (Currently Amended) A protein fraction according to claim 12, wherein liquid whey is
- 2 incorporated in to the said fraction at levels varying from 20 to 80% by weight on dry matter
- 3 basis, and that the final mixture is dried.
- 1 14. (Currently Amended) An insoluble fibre fraction produced according to [elaims 1-4] claim
- 2 1, wherein the said fraction consists of cell wall components of bran in an amount of at least 85%
- and aleurone proteins in an amount of at least 10%, and substantially free of gluten and starch,
- 4 and with a high water holding capacity of at least 6g water/g dry product.
- 1 15. (Currently Amended) A sugar fraction produced according to [claims-1-4] claim 1, wherein
- 2 the said fraction is originated primarily from the residual endosperm and it contains more than
- 3 65% sugars, such as glucose, maltose and malto-triose on dry matter basis.
- 1 16. (Currently Amended) A protein fraction derived substantially from the aleurone cells and
- 2 produced according to [claims 5-10] claim 5, wherein the said fraction contains at least 35%
- 3 protein and 10% oil, less than 5% insoluble fibre on dry matter basis, substantially free of gluten
- 4 and starch and with a high emulsifying capacity.
- 1 17. (Currently Amended) An insoluble fibre fraction produced according to [claims 5-10] claim

- 2 <u>5</u>, wherein the said fraction consists primarily of cell wall components with a relative lower
- 3 hemicellulose content compared to the original cleaned cereal bran, substantially free of gluten
- 4 and starch (<1% on dry matter basis) and with a high water holding capacity (>6g water/g dry
- 5 product).
- 1 18. (Currently Amended) A soluble hemicellulose fraction produced according to [claims 5-10]
- 2 <u>claim 5</u>, wherein the said fraction consists primarily of medium molecular weight hemicellulose
- 3 preferably above 20kDa in an amount of at least 40% of the groups arabinoxylans from wheat,
- 4 rye, rice and triticale, and beta-glucans from oat and barley, which also contains proteins in an
- 5 amount of less than 10% and monosaccharides in an amount of less than 10%, and is
- 6 substantially free of gluten and starch in an amount of less than 1% on dry matter basis.
- 1 19. (Currently Amended) A soluble oligosaccharide fraction produced according to [claims 5-
- 2 10 claim 5, wherein the said fraction consists primarily of low molecular weight hemicellulose
- 3 sub-units of below about 20kDa in an amount of at least 40% of the groups arabinoxylans from
- 4 wheat, rye, rice and triticale, and beta-glucans from oat and barley, which also contains proteins
- 5 in an amount of less than 10%, monosaccharides in an amount of less than 20%, lignans and
- 6 related phenolics in an amount of less than 5%, and is substantially free of gluten and starch in
- 7 an amount of less than 1% on dry matter basis.
- 1 20. (Currently Amended) A protein fraction according to claim 11, wherein the oil can be
- 2 optionally removed by conventional organic solvent extraction or preferably by supercritical

- 3 carbon dioxide extraction to yield an oil fraction and a defatted protein fraction.
- 1 21. (Currently Amended) A protein fraction according to claim 16, wherein the oil can be
- 2 optionally removed by conventional organic solvent extraction or preferably by supercritical
- 3 carbon dioxide extraction to yield an oil fraction and a defatted protein fraction.
- 1 22. (Currently Amended) An insoluble dietary fibre according to [any claims 14 and 17] claim
- 2 <u>14</u>, used for recovery of cellulose, hemicellulose, lignin and lignans.
- 1 23. (Currently Amended) A germ oil produced in accordance with [elaims 1-4 and 20] claim 1
- 2 containing sterols known to reduce the uptake of cholesterol in humans and intact vitamin E
- 3 complex, sterols, lecithins, phospholipids and glycolipids.
- 1 24. (Currently Amended) A defatted germ rich protein produced in accordance with [claims 1-4]
- 2 and 20] claim 1.
- 1 25. (Currently Amended) An aleurone-rich oil produced in accordance with [elaims 1-10-and
- 2 21] claim 1.
- 1 26. (Currently Amended) A defatted aleurone-rich protein produced in accordance with [elaims
- 2 1-10 and 21] claim 1.

- 1 27. (Currently Amended) A protein fraction according to [any of claims 11, 12, 13, 16, 24 and
- 2 26] claim 11, wherein proteases are incorporated in to the said fraction in wet state and at
- 3 controlled temperature and pH conditions, and the resulting protein hydrolysate has enhanced
- 4 functionalities such as solubility, emulsifying and foaming capacities.
- 1 28. (Currently Amended) The use of a protein fraction, as described in [claims 11, 12, 13, 16,
- 2 24, 26 and 27 claim 11, in feed and food applications to replace other protein products from
- 3 vegetable and animal sources.
- 1 29. (Currently Amended) The use of a protein fraction, as described in [claims 11, 12, 13, 16,
- 2 24, 26 and 27] claim 11, in food application as a texturizer, emulsifier, fat binder and fat replacer.
- 1 30. (Currently Amended) The use of a protein fraction, as described in claim 12 [and 27], as a
- 2 raw material for the extraction of soluble high-molecular weight non-starch polysaccharides.
- 1 31. (Currently Amended) The use of a protein fraction, as described in claim 12, [13 and 27] in
- 2 food applications as a foam stabilising agent, whipping agent, water binder, gelling agent, and as
- a dietary supplement rich in soluble dietary fibre (beta-glucans and arabinoxylans) with
- 4 associated health benefits such as cholesterol-reducing effects of the beta-glucans.
- 1 32. (Currently Amended) The use of a protein fraction, as described in [elaims 12, 13 and 27]
- 2 <u>claim 12</u>, as an additive or ingredient in foods such as baked products, processed meats, dairy
- 3 products, soups and sauces, high protein drinks and health drinks.

- 1 33. (Currently Amended) The use of a fibre fraction, as described in [elaims 14 and 17] claim
- 2 14, in feed and food applications to replace other insoluble fibrous products as a texturizing and
- 3 water binding additive in processed foods particularly meat products, and as a source of dietary
- 4 fibre in breakfast cereals, baked products and health products, or as a raw material for further
- 5 processing to extract remaining cellulose, hemicellulose, lignin and lignans.
- 1 34. (Currently Amended) The use of a soluble hemiceliulose, as described in claim 18, in feed
- 2 and food applications as a gellant, thickener, foam stabilizer, emulsifier, water binder, and as a
- 3 dietary supplement rich in soluble dietary fibre, and in chemical applications, or as a raw
- 4 material for further processing to obtain other functional hemicelluloses.
- 1 35. (Currently Amended) The use of a soluble hemicellulose, as described in claim 18, as an
- 2 additive or ingredient in foods such as baked products, processed meats, dairy products, soups
- and sauces, high protein drinks and health drinks.
- 1 36. (Currently Amended) The use of a soluble oligosaccharide, as described in claim 19, in feed
- 2 and food applications as a functional soluble dietary fibre or low calorie sweetener, or as a raw
- 3 material for further processing to extract lignans and associated phenolics such as ferulic acid, or
- 4 as a feedstock for industrial fermentation.
- 1 37. (Currently Amended) The use of a soluble oligosaccharide, as described in claim 19, in
- 2 confectionery formulations in combination with glucose or other sugar syrups and further
- 3 concentrated to produce moisture stable products.

- 1 38. (Currently Amended) The use of a soluble oligosaccharide, as described in claim 19, in food
- 2 and biomedical applications as a combined source of lignans and fermentable oligosaccharides
- 3 for the conversion of lignans into active cancer-reducing agents such as enterolactones.
- 1 39. (Currently Amended) The use of a sugar fraction, as described in claim 15, in .feed, food
- 2 and industrial fermentation applications as an energy source, flavouring agent and binding agent.
- 40. (Currently Amended)  $\underline{A}$  set up for carrying out the process according to [claims 1-4] claim
- 2 <u>1</u>, [characterized in that] wherein it comprises a hydrolysis vessel [(1, 8 and 11)], a wet mill [(2)],
- a heat exchange for enzymatic inactivation [(3)], decanters [(4 and 7)], a holding tank [(6)], an
- 4 ultra-filter [(10)], and optionally at least an evaporator [(13)], and dryers [(5, 9 and 12)].
- 1 41. (Currently Amended) A set up for carrying out the process according to [elaims 5-10] claim
- 2 5, [characterized in that] wherein it comprises a hydrolysis vessels [(1, 8 and 11)], a wet mill
- 3 [(2)], a heat exchange for enzymatic inactivation [(3)], decanters [(4 and 7)], a holding tank [(6)],
- 4 an ultra-filter [(10)], and optionally evaporators [(12 and 13)], and dryers [(5 and 9)].
- 1 42. (Currently Amended) A process according to [elaims 1-4] claim 1, wherein the enzymatic
- 2 treatment is carried out for less than 3 hours at a pH of 4 to 7.5 and at a temperature of from 50
- 3 to 90°C, at an enzymatic activity of at least 1 IU/g of substrate, preferably 200 to 1500 IU/g of
- 4 substrate.

- 1 43. (Currently Amended) A process according to [elaims 5-10] claim 5, wherein the enzymatic
- 2 treatment is carried out for less than 3 hours at a pH of 4 to 7, preferably 4.5-5.5, and at a
- 3 temperature of from 35 to 80°C, at an enzymatic activity of at least 1 IUlg of substrate,
- 4 preferably 200 to 1500 IU/g of substrate.